CHILDRENS ALLERGY- diphenhydramine hydrochloride liquid Chain Drug Consortium, LLC

Active ingredient

Drug Facts

Active ingredient (in each 5 mL teaspoon)

Diphenhydramine HCL 12.5 mg

Purpose

Purpose

Antihistamine

Keep out of reach of children

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away.

Uses

Uses

- Temporarily relieves these symptoms due to hay fever or other respiratory allergies:
 - sneezing
 - itching of the nose or throat
 - runny nose
 - itchy watery eyes
- temporarily relieves these symptoms due to the common cold:
 - sneezing
 - runny nose

Warnings

Warnings

Do not use

- with any other product containing diphenhydramine, even one used on skin
- to make a child sleepy

Ask a doctor before use Ask a doctor or pharmacist before use Ask a doctor before use if the child has

- glaucoma
- a breathing problem such as chronic bronchitis

Ask a doctor or pharmacist before if the child is

taking sedatives or tranquilizers

When using this product When using this product

- marked drowsiness may occur
- excitability may occur, especially in children
- · sedatives and tranquilizers may increase drowsiness

Directions

Directions

- if needed, take every 4-6 hours
- do not take more than 6 doses in 24 hours

Children under 4 years of age: do not use

Children 4 to under 6 years of age: do not use unless directed by

a doctor

Children 6 to under 12 years of age: 1 to 2 teaspoonfuls (12.5 mg

to 25 mg)

Other Information

Other Information

- Keep carton for full directions for use
- each teaspoonful contains: sodium 10 mg
- store at 20-25 ° C (68-77 ° F)
- dosage cup provided

Inactive ingredients

Inactive ingredients

Citric acid, D and C Red # 40, flavor, glycerin, poloxamer 407, purified water, sodium benzoate, sodium chloride, sodium citrate, sucrose

Questions or comments?

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Call weekdays from 9:30 AM to 4:30 PM EST at

1-877-798-5944

Product Label

NDC 68016-0823-54

*COMPARE TO THE ACTIVE INGREDIENT IN BENADRYL® ALLERGY LIQUID

PREMIER VALUE ®

CHILDREN'S ALLERGY

DIPHENHYDRAMINE HYDROCHLORIDE ANTIHISTAMINE

Relieves: Sneezing, Runny Nose, Itchy Watery Eyes, Itchy Throat

Alcohol-Free Cherry Flavor

4 FL OZ (118 mL)

INDEPENDENTLY TESTED SATISFACTION GUARANTEED PV

If for any reason you are not satisfied with this product, please return it to the store where purchased for a full refund.

DISTRIBUTED BY: CHAIN DRUG CONSORTIUM 3301 NW BOCA RATON BLVD SUITE 101, BOCA RATON, FL 33431

BX-005

DO NOT USE IF PRINTED SEAL UNDER CAP IS TORN OR MISSING

*This product is not manufactured or distributed by McNeil-PPC, Inc. distributer of Benadryl® Allergy Liquid.

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6 years of age:	directed by a doctor
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12 years of age:	(12.5 mg to 25 mg)

Drug Facts (continued)

Other Information

- Keep carton for full directions for use
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- store at 20° 25° C (68° 77° F)
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NOC 88016-0123-54 "COMPARE TO THE ACTIVE INGREDIENT IN BENADRYL® ALLERSY LIQUID



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BX-005

Relieves: Sneezing, Runny Nose, Itchy Watery Eyes, **Itchy Throat**

Alcohol-Free Cherry Flavor

4 FL OZ (118 mL)



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Drug Facts (continued)

Other Information

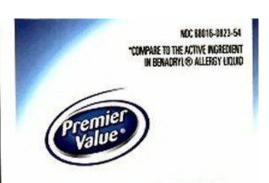
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BX-005

Relieves: Sneezing, Runny Nose, Itchy Watery Eyes, **Itchy Throat**

Alcohol-Free Cherry Flavor

4 FL OZ (118 mL)



CHILDRENS ALLERGY

diphenhydramine hydrochloride liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:68016-823

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength **DIPHENHYDRAMINE HYDROCHLORIDE** (UNII: TC2D6|AD40) **DIPHENHYDRAMINE** 12.5 mg (DIPHENHYDRAMINE - UNII:8GTS82S83M) **HYDROCHLORIDE** in 5 mL

Inactive Ingredients

Ingredient Name

Strength

CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SUCROSE (UNII: C151H8M554)	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:68016-823-	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/11/2012	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M009	09/11/2012	

Labeler - Chain Drug Consortium, LLC (101668460)

Registrant - AptaPharma Inc. (790523323)

Establishment			
Name	Address	ID/FEI	Business Operations
AptaPharma Inc.		790523323	manufacture(68016-823)

Revised: 12/2023 Chain Drug Consortium, LLC